## Claims:

We claim:

1. A method of killing or inhibiting a microorganism, comprising contacting said microorganism with a composition comprising:

- (a) a peroxidase produced by or derived from the fungus Coprinus;
  - (b) an enhancing agent; and
  - (c) a hydrogen peroxide or a source of hydrogen peroxide.
- 10 2. The method of claim 1, wherein the peroxidase is a recombinant enzyme obtainable from *Coprinus cinereus*.
  - 3. The method of claim 1; wherein the peroxidase is obtainable from *Coprinus cinereus*, IFO 8371.
  - 4. The method of claim 1, wherein the source of hydrogen peroxide is an enzymatic hydrogen peroxide-generating system.
- 5. The method of claim 4, wherein the enzymatic system is selected from the group consisting of glucose oxidase/glucose, hexose oxidase/hexose, L- or D-amino acid oxidase/L- or D-amino acid, and lactate oxidase/lactate.
- 6. The method of claim 1, wherein the enhancing agent is an 25 electron donor.
  - 7. The method of claim 1; wherein the enhancing agent is a water-soluble halide or thiocyanate salt.
- 30 8. The method of claim 1; wherein the enhancing agent is a compound having the formula:

in which formula X represents (-O-) or (-S-), and the 10 substituent groups R<sup>1</sup>-R<sup>9</sup>, which may be identical or different, independently represents any of the following radicals: hydrogen, halogen, hydroxy, formyl, carboxy, and esters and salts hereof, carbamoyl, sulfo, and esters and salts hereof, sulfamoyl, nitro, amino, phenyl, C1-C14-alkyl, C1-C5-alkoxy, 15 carbonyl-C<sub>1</sub>-C<sub>5</sub>-alkyl, aryl-C<sub>1</sub>-C<sub>5</sub>-alkyl; which carbamoyl, sulfamoyl, and amino groups may be unsubstituted or substituted once or twice with a substituent group R<sup>10</sup>; and which phenyl may be unsubstituted or substituted with one or more substituent groups  $R^{10}$ ; and which  $C_1-C_{14}$ -alkyl,  $C_1-C_5$ -alkoxy, carbonyl- $C_1-C_5$ -20 alkyl, and aryl-C<sub>1</sub>-C<sub>5</sub>-alkyl groups may be saturated or unsaturated, branched or unbranched, and may be unsubstituted or substituted with one or more substituent groups R10; which substituent group R10 represents any of the following radicals: halogen, hydroxy, formyl, carboxy and esters or salts 25 thereof, carbamoyl, sulfo and esters or salts thereof, sulfamoyl, nitro, amino, phenyl, aminoalkyl, piperidino, piperazinyl, pyrrolidin-1-yl, C<sub>1</sub>-C<sub>5</sub>-alkyl, C<sub>1</sub>-C<sub>5</sub>-alkoxy; which carbamoyl, sulfamoyl, and amino groups may be unsubstituted or substituted once or twice with hydroxy, C<sub>1</sub>-C<sub>5</sub>-alkyl, or C<sub>1</sub>-C<sub>5</sub>-30 alkoxy; and which phenyl may be substituted with one or more of the following radicals: halogen, hydroxy, amino, formyl, carboxy

and esters and salts hereof, carbamoyl, sulfo and esters and

acid.

salts hereof, and sulfamoyl; and which C<sub>1</sub>-C<sub>5</sub>-alkyl, and C<sub>1</sub>-C<sub>5</sub>-alkoxy groups may be saturated or unsaturated, branched or unbranched, and may be substituted once or twice with any of the following radicals: halogen, hydroxy, amino, formyl, carboxy and seters and salts thereof, carbamoyl, sulfo and esters and salts hereof, and sulfamoyl; or in which general formula two of the substituent groups R<sup>1</sup>-R<sup>9</sup> may together form a group -B-, in which B represents any of the following the groups: (-CHR<sup>10</sup>-N=N-), (-CH=CH-)<sub>n</sub>, (-CH=N-)<sub>n</sub> or (-N=CR<sup>10</sup>-NR<sup>11</sup>-), in which groups n-represents an integer of from 1 to 3, R<sup>10</sup> is a substituent group as defined above and R<sup>11</sup> is defined as R<sup>10</sup>.

- 9. The method of claim 1, wherein the enhancing agent is 15 selected from the group consisting of 10-methylphenothiazine, phenothiazine-10-propionic acid, N-hydroxysuccinimide phenothiazine-10-propionate, 10-ethyl-phenothiazine-4-carboxylic acid, 10-ethylphenothiazine, 10-propylphenothiazine, 10isopropylphenothiazine, methyl phenothiazine-10-propionate, 10-20 phenylphenothiazine, 10-allylphenothiazine, 10-(3-(4methylpiperazin-1-yl)propyl)phenothiazine, 10-(2-pyrrolidin-1yl-ethyl) phenothiazine, 2-methoxy-10-methyl-phenothiazine, 1methoxy-10-methylphenothiazine, 3-methoxy-10methylphenothiazine, 3,10-dimethylphenothiazine, 3,7,10-25 trimethylphenothiazine, 10-(2-hydroxyethyl)phenothiazine, 10-(3hydroxypropyl) phenothiazine, 3-(2-hydroxyethyl)-10methylphenothiazine, 3-hydroxymethyl-10-methylphenothiazine, 3,7-dibromophenothiazine-10-propionic acid, phenothiazine-10propionamide, chlorpromazine, 2-chloro-10-methylphenothiazine, 30 2-acetyl-10-methylphenothiazine, 10-methylphenoxazine, 10-ethylphenoxazine, phenoxazine-10-propionic acid, 10-(2hydroxyethyl)phenoxazine and 4-carboxyphenoxazine-10-propionic
- 35 10. The method of claim 1; wherein the enhancing agent is a compound having the formula:

wherein A denotes a group -D, -CH=CH-D, -CH=CH-CH=CH-D, -CH=N-D, -N=N-D, or -N=CH-D; D is selected from the group consisting of -CO-E, -SO<sub>2</sub>-E, - N-XY, and -N<sup>+</sup>-XYZ; E is -H, -OH, -R, or 5 -OR, and X and Y and Z may be identical or different and selected from -H and -R; R is  $C_1$ - $C_{16}$  alkyl, preferably saturated or unsaturated, branched or unbranched  $C_1$ - $C_8$  alkyl, optionally substituted with a carboxy, sulfo or amino group; and B and C may be the same or different and selected from  $C_mH_{2m+1}$ ;  $1 \le m \le 10.5$ .

- 11. The method of claim 1, wherein the enhancing agent is acetosyringone, methylsyringate, ethylsyringate, propylsyringate, butylsyringate, hexylsyringate, or octylsyringate.
- 12. The method of claim 1, wherein the enhancing agent is a salt selected from the group consisting of potassium halide, sodium halide, lithium halide, ammonium halide, calcium halide.
- 20 13. The method of claim 1, wherein the enhancing agent is a salt salt selected from the group consisting of potassium iodide, sodium iodide, lithium iodide, ammonium iodide, and calcium iodide.
- 25 14. The method of claim 1, wherein the enhancing agent is a salt selected from the group consisting of sodium thiocyanate, potassium thiocyanate, and ammonium thiocyanate
- 15. The method of claim 1, wherein the microorganism is present 30 in laundry.

- 16. The method of claim 1, wherein the microorganish is present on skin, hair, mucous membranes, teeth, wounds, bruises or in the eye or oral cavity, of a human or animal.
- 5 17. The method of claim 1, wherein the composition is in the form of a soaking, washing or rinsing liquor.
  - 18. The method of claim 1, wherein the composition is a liquid composition.
- 19. The method of claim 1, wherein the composition is a mouth wash, an antiinflammatory liquid, a perspirant, a deodorant, or a nasal spray.
- 15 20. The method of claim 1, wherein the composition is a solid composition.
- 21. The method of claim 1, wherein the composition is an eye ointment, an anti-inflammatory ointment, a foot bath salt, a 20 perspirant, or a deodorant.
  - 22. A method of killing or inhibiting a microorganism, comprising contacting said microorganim with a composition comprising:
- (a) a peroxidase;
  - (b) an enhancing agent of the formula:

wherein A denotes a group -D, -CH=CH-D, -CH=CH-CH=CH-D, -CH=N-D, -N=N-D, or -N=CH-D; D is selected from the group consisting of -CO-E, -SO<sub>2</sub>-E, -N-XY, and -N<sup>+</sup>-XYZ; E is -H, -OH, -R, or -OR, and X and Y and Z may be identical or different and selected from -H and -R; R is  $C_1$ - $C_{16}$  alkyl, preferably saturated or unsaturated, branched or unbranched  $C_1$ - $C_8$  alkyl, optionally substituted with a carboxy, sulfo or amino group; and B and C may be the same or different and selected from  $C_mH_{2m+1}$ ;  $1 \le m \le 5$ ; and

- (c) a hydrogen peroxide or a source of hydrogen peroxide.
  - 23. A method of preserving a cosmetic product, comprising adding to the cosmetic product an effective amount of an enzymatic antimicrobial composition comprising:
- (a) a peroxidase produced by or derivable from the fungus Coprinus;
  - (b) an enhancing agent; and
  - (c) hydrogen peroxide or a source of hydrogen peroxide.
- 20 24. The method according to claim 23, wherein the cosmetic product is a liquid, a gel, a paste, an ointment or a lotion.
- 25. The method according to claim 23, wherein the cosmetic product is a mouth wash, an eye lotion, a perspirant, a 25 deodorant, a nasal spray, an eye ointment, or a foot bath salt.
  - 26. A method for cleaning or disinfecting contact lenses comprising contacting said contact lenses with an effective amount of an a enzymatic antimicrobial composition comprising:
- (a) a peroxidase produced by or derivable from the fungus Coprinus;
  - (b) an enhancing agent; and
  - (c) hydrogen peroxide or a source of hydrogen peroxide.
- 35 27. A method of inhibiting microbial growth on a hard surface, wherein the surface is contacted with an a enzymatic antimicrobial composition comprising:

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- (a) a peroxidase produced by or derivable from the fungus Coprinus;
  - (b) an enhancing agent; and
  - (c) hydrogen peroxide or a source of hydrogen peroxide.
- 28. The method according to claim 27, wherein the hard surface is a process equipment member of a cooling tower, a water treatment plant, a dairy, a food processing plant, a chemical process plant or pharmaceutical process plant.
- 29. The method according to claim 27, wherein the hard surface is a surface of water sanitation equipment.
- 30. The method according to claim 27, wherein the hard surface 15 is a surface of equipment for paper pulp processing.